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(71) Applicant (*for all designated States except US*): SMITHKLINE BEECHAM CORPORATION [US/US]; One Franklin Plaza, P.O. Box 7929, Philadelphia, Pennsylvania 19101 (US).

(72) Inventors; and

(75) Inventors/Applicants (*for US only*): FAUCHER, Nicolas Eric [FR/FR]; GlaxoSmithKline, Centre de Recherches, ZA de Courtaboeuf, 25 Avenue du Quebec, F-91940 Les Ulis (FR). MARTRES, Paul [FR/FR]; GlaxoSmithKline, Centre de Recherches, ZA de Courtaboeuf, 25 Avenue du Quebec, F-91940 Les Ulis (FR).

(74) Agent: LEAROYD, Stephanie, Anne; GlaxoSmithKline, Corporate Intellectual Property (CN925.1), 980 Great West Road, Brentford, Middlesex TW8 9GS (GB).

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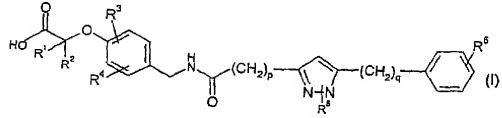
(54) Title: SUBSTITUTED PYRAZOLES AS PPAR AGONISTS



(57) Abstract: A compound of formula (I) and pharmaceutically acceptable salts, solvates and hydrolysable esters thereof (I) wherein: p is O or 1; q is O or 1; R^1 and R^2 are independently H or C_{1-3} alkyl; R^3 and R^4 are independently H, C_{1-6} alkyl, $-OC_{1-6}$ alkyl, halogen, OH, C_{2-6} alkenyl or CF_3; R^5 is H, C_{1-6} alkyl (optionally substituted by one or more halogens, -COphenyl, OC_{1-6} alkyl, phenyl morpholino or C_{2-6} alkenyl. R^6 is C_{1-6} alkyl, halogen, $-OCH_2$ phenyl, phenyl (optionally substituted by C_{1-3} alkyl), morpholino, pyrrolidino, piperidino, thiophenyl, furanyl pyridinyl or $-OC_{2-6}$ alkenyl. These compounds activate the alpha and gamma subtypes fo the hppar receptor and are useful e.g. in the treatment of diabetes, dyslipidemia or syndrome X.

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